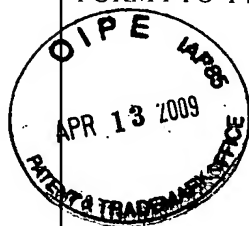


FORM PTO-1449

U. S. DEPARTMENT OF COMMERCE  
PATENT AND TRADEMARK OFFICEINFORMATION DISCLOSURE  
STATEMENT BY APPLICANT

(Use several sheets if necessary)

ATTY. DOCKET NO.

U 015859-4

SERIAL NO.

10/542,937

APPLICANT

PEDRO MATA LOPEZ, ET AL.

FILING DATE

SEPTEMBER 8, 2006

GROUP

1634

## U.S. PATENT DOCUMENTS

EXAMINER INITIALS	REFERENCE DESIGNATION	DOCUMENT NUMBER	DATE	NAME	FILING DATE IF APPROPRIATE
	AA				
	AB				
	AC				
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## FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	TRANSLATION	
					YES	NO
	AL					
	AM					
	AN					
	AO					
	AP					

## OTHER ART (Including Author, Title, Date, Pertinent Dates, Etc.)

	AR	Tejedor, Diego, et al. "Reliable Low-Density DNA Array Based on Allele-Specific Probes for Detection of 118 Mutations Causing Familial Hypercholesterolemia", Clinical Chemistry Vol. 51:7 pages 1137-1144 (2005)
	AS	Alonso, R. "Cardiovascular disease in familial hypercholesterolemia: Influence of low-density lipoprotein receptor mutation type and classic risk factors", Atherosclerosis, pages 1-7 (2008)
	AT	Civerira, Fernando, et al. "Comparison of Genetic Versus Clinical Diagnosis in Familial Hypercholesterolemia" The American Journal of Cardiology(www.AJConline.org), pages 1-7 (2008).

EXAMINER

DATE CONSIDERED

EXAMINER:

Initial if citation considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

<b>FORM PTO-1449 U. S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE</b>  <b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  (Use several sheets if necessary)		<b>ATTY. DOCKET NO.</b>		<b>SERIAL NO.</b>	
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		<b>APPLICANT</b>			
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	AP					

OTHER ART (Including Author, Title, Date, Pertinent Dates, Etc.)		
	AR	Junyent, Mireia, et al. "Femoral Atherosclerosis in Heterozygous Familial Hypercholesterolemia: Influence of the Genetic Defect", Arterioscler Thromb Vasc Biol Vol. 28, pages 580-586 (2008)
	AS	Mozas Pilar, et al. "Molecular Characterization of Familial Hypercholesterolemia in Spain: Identification of 39 Novel and 77 Recurrent Mutations in LDLR", Human Mutation Mutation in Brief Vol. 735 Pages 1-13 (2004)
	AT	

<b>EXAMINER</b>	<b>DATE CONSIDERED</b>
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